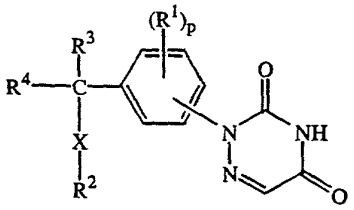
ABSTRACT

IL-5 INHIBITING 6-AZAURACIL DERIVATIVES

The present invention is concerned with the compounds of formula



(I)

the *N*-oxides, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, wherein p is 0 to 4; X is O, S, NR^5 or a direct bond; Y is O, S, NR^5 or $S(O)_2$; R^1 independently is C_{1-6}alkyl, halo, polyhaloC_{1-6}alkyl, hydroxy, mercapto, C_{1-6}alkyloxy, C_{1-6}alkylthio, C_{1-6}alkylcarbonyloxy, aryl, cyano, nitro, Het^3, R^6, NR^7R^8 or substituted C_{1-4}alkyl; R^2 is Het^1, C_{3-7}cycloalkyl or optionally substituted C_{1-6}alkyl and if X is O, S or NR^5, then R^2 may also represent aminocarbonyl, aminothiocarbonyl, C_{1-4}alkylcarbonyl, C_{1-4}alkylthiocarbonyl, arylcarbonyl, arylthiocarbonyl, Het^1carbonyl or Het^1thiocarbonyl; R^3 and R^4 independently are hydrogen, C_{1-6}alkyl or C_{3-7}cycloalkyl; R^3 and R^4 form a C_{2-6}alkanediyl; R^5 is hydrogen or C_{1-4}alkyl; R^6 is a sulfonyl or sulfinyl derivative; R^7 and R^8 are independently hydrogen, optionally substituted C_{1-4}alkyl, aryl, a carbonyl containing moiety, C_{3-7}cycloalkyl, -Y-C_{1-4}alkanediyl-C(=O)-O-R^{14}, Het^3, Het^4 and R^6; R^{11} is hydroxy, mercapto, cyano, nitro, halo, trihalomethyl, C_{1-4}alkyloxy, formyl, trihaloC_{1-4}alkylsulfonyloxy, R^6, NR^7R^8, C(=O)NR^7R^8, C_{1-4}alkanediyl-C(=O)-O-R^{14}, -C(=O)-O-R^{14}, -Y-C_{1-4}alkanediyl-C(=O)-O-R^{14}, aryl, aryloxy, arylcarbonyl, C_{3-7}cycloalkyl, C_{3-7}cycloalkyloxy, phthalimide-2-yl, Het^3 and C(=O)Het^3; R^{14} is hydrogen, C_{1-4}alkyl, C_{3-7}cycloalkyl, aminocarbonylmethylene or mono-or di(C_{1-4}alkyl)aminocarbonylmethylene; aryl is optionally substituted phenyl; Het^1, Het^2, Het^3 and Het^4 are optionally substituted heterocycles; to processes for their preparation and compositions comprising them. It further relates to their use as a medicine.